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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

First Named Inventor: Stenkamp, Dirk, et al) Art Unit: 1626

Application No.: 10/699,089) Examiner: Powers, Fiona

Filing Date: October 31, 2003) Confirmation No.: 1853

Title: Amide Compounds Having MCH-Antagonistic Activity and Medicaments Comprising these Compounds

Attorney Docket No.: 1/1407

IDS STATEMENT(S)

INFORMATION PREVIOUSLY CITED OR SUBMITTED

U.S. Application No. ("the parent application") is relied on by the above-identified application for an earlier effective filing date under 35 U.S.C. 120. The patents, publications, pending U.S. applications, or other information listed in the instant information disclosure statement were previously cited by the Office in the parent application or submitted to the Office in the parent application in an information disclosure statement that complied with 37 CFR 1.98(a)-(c). 37 CFR 1.98(d)

The fee set forth in 37 CFR 1.17(p) has been submitted herewith. 37 CFR 1.97(c-d) Otherwise, the Applicant authorizes the charging of all fees set forth in §1.16 and to §1.17 to Deposit Account No 02-29955 for the entire pendency of this application. 37 CFR 1.25(b)

INFORMATION FIRST CITED BY FOREIGN PATENT OFFICE NOT MORE THAN THREE (3) MONTH PRIOR

Each item of information contained in the instant information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three (3) months prior to the filing of the instant information disclosure statement. 37 CFR 1.97(e)(1)

NO INFORMATION KNOWN MORE THAN THREE (3) MONTHS PRIOR

No item of information contained in the instant information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing this certification after making reasonable inquiry, no item of information contained in the instant information disclosure statement was known to any individual designated in 37 C.F.R. §1.56(c) more than three (3) months prior to the filing of the instant information disclosure statement. 37 CFR 1.97(e)(2)

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37 CFR 1.704(d)

Respectfully submitted,



Timothy X. Witkowski

Reg. No. 40,232

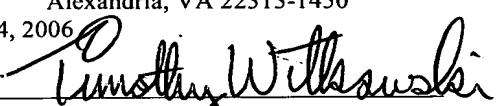
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Timothy X. Witkowski 40,232



PTO/SB/08a (08-03)
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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(Use as many sheets as necessary)

Sheet

1 of 10

Complete if Known

Application Number	10/699,089
Filing Date	October 31, 2003
First Named Inventor	Stenkamp, D., et al
Art Unit	1614
Examiner Name	To be assigned

Attorney Docket Number 1/1407

U. S. PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	Document Number Number-Kind Code ² (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		US- 3,994, 900	11/30/1976	Krapcho, John, et al	
		US- 6,366,268 B1	04/02/2002	Forrest, S.R., et al	
		US- 2002/0052383 A1	05/02/2002	Bakthavatchalam, R. et al	
		US- 2005/0026915 A1	02/03/2005	DeVita, R.J., et al	
		US- 2005/0009815 A1	01/13/2005	DeVita, R.J., et al	
		US- 2004/0082780 A1	04/29/2004	Doherty, E.M., et al	
		US- 2003/0192132 A1	10/16/2003	Chassot, L., et al	
		US- 2004/0220191 A1	11/04/2004	Schwink, L., et al	
		US- 2004/0142953 A1	07/22/2004	Delorme, D., et al	
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		US-			

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	Foreign Patent Document Country Code ³ -Number ⁴ Kind Code ⁵ (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ⁶
		WO 2005/063239 A1	07/14/2005	Boehringer Ingelheim International GmbH		
		WO 2005/085221 A1	09/15/2005	Boehringer Ingelheim International GmbH		
		DE 1 088 955	09/15/1960	Farbenfabriken Bayer Ak		
		DE 3016827	11/20/1980	Continental Pharam S.A., Brussel		
		WO 2004/024702	03/25/2004	Boehringer Ingelheim Pharma GmbH		

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		Country Code ³ Number ⁴ Kind Code ⁵ (if known)			
		EP 1 593 667 A1	09/11/2005	Takeda Pharm. Co.	
		WO 2004/018467 A2	03/04/2004	BI Pharma GmbH	
		WO 2004/041820 A1	05/21/2004	BI Pharma GmbH	
		WO 03/045313 A2	06/05/2003	Merck & Co. Inc.	
		EP 1 283 199 A1	02/12/2003	Takeda Chem. Ind.	

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		Country Code ³ Number ⁴ Kind Code ⁵ (if known)				
	WO 03/014111 A1		02/20/2003	Astrazeneca Ab		
	WO 02/28182 A1		04/11/2002	Syngenta Participation , Ag		
	WO 03/050087 A2		06/19/2003	Syngenta Participation Ag		
	WO 98/38156		09/03/1998	Takeda Chemical, Inc.		
	WO 03/024448 A2		03/27/2003	Methylgene, Inc.		

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		Country Code ³ ·Number ⁴ ·Kind Code ⁵ (if known)				
		EP 0 237 678 A1	09/23/1987	Glaxo Group Ltd.		
		WO 99/29674	06/17/1999	Janssen Pharmaceuticals		
		JP 04054118	02/21/1992	Tanabe Seiyaku Co.		
		WO 2004/072018	08/26/2004	Takeda Chemicals, Inc.		
		WO 2000/05223	02/02/2000	Zeneca Ltd., Inc.		

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Sheet	6	of	10	Attorney Docket Number	
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		Country Code ³ Number ⁴ Kind Code ⁵ (if known)			
	WO 2000/49005		08/24/2000	Aventis Pharmaceuticals, Ltd.	
	WO 2001/02344		01/11/2001	Taisho Pharmaceuticals,	
	JP 2000/086603		03/28/2000	Yoshitomi Pharmaceuticals,	
	WO 02/051809 A1		07/04/2002	Schering Corporation	
	WO 03/032980 A1		04/24/2003	Glaxo Group Ltd.	

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Attorney Docket Number

Complete if Known

10/699 089

October 31, 2003

Stenkamp D. et al

1614

To be assigned

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		Country Code ³ Number ⁴ Kind Code ⁵ (if known)				
	WO 03/033476 A1		04/24/2003	SmithKline Beecham Plc.		
	WO 99/02497 A2		01/21/1999	Novartis-Erfindungen		
	WO 02/092068		11/21/2002	Ono Pharmaceuticals		
	WO 03/045920 A1		06/05/2003	Merck & Co., Inc.		
	WO 03/049702		06/19/2003	Amgem, Inc.		

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		Country Code ³ Number ⁴ Kind Code ⁵ (if known)			
		EP 0 810 220 A1	12/03/1997	Pfizer, Inc.	
		WO 02/079144 A1	10/10/2002	Wella Aktien.	
		WO 2004/072025 A2	08/26/2004	Aventis Pharmaceuticals ,	
		WO 2004/069823 A1	08/19/2004	Methylgene, Inc.	
		WO 2004/037751 A2	05/06/2004	Temple University	

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Sheet	9	of	10	Attorney Docket Number	1/1407

NON PATENT LITERATURE DOCUMENTS				
Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.		T ²
		YANYUN CHEN, et al. "Targeted Disruption of the Melanin-Concentrating Hormone Receptor-1 Results in Hyperphagia and Resistance to Diet-Induced Obesity", Endocrinology 143(7):2469-2477 2002		
		DAQING QU, et al. "A role for melanin-concentrating hormone in the central regulation of feeding behaviour" Nature Vol. 380, pp 243-247 1996		
		MASAKO SHIMADA, et al. "Mice lacking melanin-concentrating hormone are hypophagic and lean" Nature Vol. 396, pp 670-674 1998		
		BETH BOROWSKY, et al. "Antidepressant, anxiolytic and anorectic effects of a melanin-concentrating hormone-1 receptor antagonist" Nature Medicine Vol. 8, No. 8, pp 825-830, 2002		
		DONALD J. MARSH, et al. "Melanin-concentrating hormone 1 receptor-deficient mice are lean, hyperactive, and hyperphagic and have altered metabolism" PNAS, Vol. 99, No. 5, pp 3240-3245, 2002		
		SHIRO TAKEKAWA, et al. "T-226296: a novel, orally active and selective melanin-concentrating hormone receptor antagonist" E. Journal of Pharm. Vol. 438, pp 129-13, 2002		
		J. KRAPCHO, et al; "Immunosuppressive Activity of 2'-(3-Dimethylaminopropylthio)cinnamanilide (Cinanserin) and Related Compounds" J. Med. Chemistry. 1969, 12(1), 164-166		
		TAMIZ, Amir P., et al; Structure-Activity Relationship of N-(Phenylalkyl)cinnamides as Novel NR2B Subtype-Selective NMDA Receptor Antagonists - J. Med. Chem. 1999, 42, 3412-3420		
		TETSUO, Sato, et al; Japanese Abstract 2002193800; VEGF Receptor antagonists for Treatment of neoangiogenesis-related diseases		

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1 Applicant's unique citation designation number (optional). 2 Applicant is to place a check mark here if English language Translation is attached. This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: **Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.**

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Substitute for form 1449B/PTO				<i>Complete if Known</i>	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT				Application Number	10/699,089
				Filing Date	October 31, 2003
				First Named Inventor	Stenkamp, D., et al
				Art Unit	1614
				Examiner Name	To be assigned
(Use as many sheets as necessary)				Attorney Docket Number	
Sheet	10	of	10	1/1407	

NON PATENT LITERATURE DOCUMENTS

Examiner Signature		Date Considered	
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PTO/SB/17 (01-06)

Approved for use through 07/31/2006. OMB 0651-0032
U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

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Fees pursuant to the Consolidated Appropriations Act, 2005 (H.R. 4818).

FEE TRANSMITTAL For FY 2006

 Applicant claims small entity status. See 37 CFR 1.27

TOTAL AMOUNT OF PAYMENT (\$)

Complete if Known

Application Number	10/699,089
Filing Date	October 31, 2003
First Named Inventor	Stenkamp, Dirk, et al
Examiner Name	Powers, Fiona
Art Unit	1626
Attorney Docket No.	1/1407

METHOD OF PAYMENT (check all that apply)

Check Credit Card Money Order None Other (please identify): _____
 Deposit Account Deposit Account Number: 02-2955 Deposit Account Name: Boehringer Ingelheim Corporation
 For the above-identified deposit account, the Director is hereby authorized to: (check all that apply)
 Charge fee(s) indicated below Charge fee(s) indicated below, except for the filing fee
 Charge any additional fee(s) or underpayments of fee(s) Credit any overpayments
 under 37 CFR 1.16 and 1.17

WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.

FEE CALCULATION (All the fees below are due upon filing or may be subject to a surcharge.)

1. BASIC FILING, SEARCH, AND EXAMINATION FEES

Application Type	FILING FEES		SEARCH FEES		EXAMINATION FEES	
	Fee (\$)	Small Entity	Fee (\$)	Small Entity	Fee (\$)	Small Entity
Utility	300	150	500	250	200	100
Design	200	100	100	50	130	65
Plant	200	100	300	150	160	80
Reissue	300	150	500	250	600	300
Provisional	200	100	0	0	0	0

2. EXCESS CLAIM FEES

Fee Description

Each claim over 20 (including Reissues) Fee (\$)
 Each independent claim over 3 (including Reissues) Fee (\$)
 Multiple dependent claims Fee (\$)

Total Claims	Extra Claims	Fee (\$)	Fee Paid (\$)	Multiple Dependent Claims	
				Fee (\$)	Fee (\$)
- 20 or HP =	x	=		50	25

HP = highest number of total claims paid for, if greater than 20.

Indep. Claims	Extra Claims	Fee (\$)	Fee Paid (\$)	Multiple Dependent Claims	
				Fee (\$)	Fee Paid (\$)
- 3 or HP =	x	=			

HP = highest number of independent claims paid for, if greater than 3.

3. APPLICATION SIZE FEE

If the specification and drawings exceed 100 sheets of paper (excluding electronically filed sequence or computer listings under 37 CFR 1.52(e)), the application size fee due is \$250 (\$125 for small entity) for each additional 50 sheets or fraction thereof. See 35 U.S.C. 41(a)(1)(G) and 37 CFR 1.16(s).

Total Sheets	Extra Sheets	Number of each additional 50 or fraction thereof	Fee (\$)	Fee Paid (\$)
- 100 =	/ 50 =	(round up to a whole number) x	=	

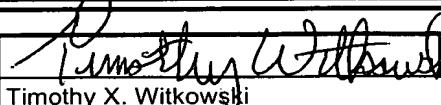
4. OTHER FEE(S)

Non-English Specification, \$130 fee (no small entity discount)

Other (e.g., late filing surcharge): Information Disclosure Statement Fees Paid (\$)

180.00

SUBMITTED BY

Signature		Registration No. 40,232 (Attorney/Agent)	Telephone 203-798-4310
Name (Print/Type)	Timothy X. Witkowski		Date May 24, 2006

This collection of information is required by 37 CFR 1.136. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 30 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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